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OM protein - protein search, using sw model

Run on: January 7, 2002, 15:40:13 ; Search time 154.28 Seconds
(without alignments)
22.086 Million cell updates/sec

Title: US-08-569-749-8

Perfect score: 267
Sequence: 1 LAKAGFYIGFGDRVACFPAC.....WPKDNMSHLRHPKCPF 46

Scoring table:

BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 522463 seqs, 74073290 residues

Total number of hits satisfying chosen parameters: 522463

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

Database :

A_Geneseq_1101:*

1: /SID52/gcgdata/geneseq/geneseqp/AA1980.DAT:*
2: /SID52/gcgdata/geneseq/geneseqp/AA1981.DAT:*
3: /SID52/gcgdata/geneseq/geneseqp/AA1982.DAT:*
4: /SID52/gcgdata/geneseq/geneseqp/AA1983.DAT:*
5: /SID52/gcgdata/geneseq/geneseqp/AA1984.DAT:*
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8: /SID52/gcgdata/geneseq/geneseqp/AA1987.DAT:*
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20: /SID52/gcgdata/geneseq/geneseqp/AA1999.DAT:*
21: /SID52/gcgdata/geneseq/geneseqp/AA2000.DAT:*
22: /SID52/gcgdata/geneseq/geneseqp/AA2001.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	267	100.0	46	AAW13550	Human c-IAP2 repa
2	267	100.0	604	AAW19747	Human inhibitor of
3	267	100.0	604	AAW13546	Human c-IAP2. Hom
4	267	100.0	604	AAW52703	Human cellular inh
5	267	100.0	604	AAW33997	Human cellular inh
6	267	100.0	1141	AAW50694	Human API2-MIT chi
7	264	98.9	604	AAW19582	Human apoptosis in
8	264	98.9	604	AAW69295	Human HIAP-1 prote
9	259	97.0	306	AAU02935	Angiotensin conver
10	248	92.9	46	AAW13548	Human c-IAP1 repa
11	248	92.9	438	AAW04583	Human inhibitor of

12	248	92.9	618	AAW19746	Human inhibitor of
13	248	92.9	618	AAW19583	Human apoptosis in
14	248	92.9	618	AAW13545	Human c-IAP1. Hom
15	248	92.9	618	AAW69296	Human HIAP-2 prote
16	248	92.9	618	AAW33998	Human cellular inh
17	247	92.5	612	AAW13555	Human c-IAP. Mus
18	247	92.5	612	AAW69299	Murine HIAP-2 prot
19	244	91.4	591	AAW19586	Murine apoptosis in
20	235	88.0	600	AAW59298	Mouse apoptosis in
21	235	88.0	602	AAW19285	Murine HIAP-1 prot
22	182	68.2	497	AAW19581	Mouse apoptosis in
23	182	68.2	497	AAW69294	Human XIAP protein
24	182	68.2	497	AAW39985	Human X-linked inh
25	182	68.2	497	AAW39981	Human XIAP protein
26	177	66.3	496	AAW19745	Mouse inhibitor of
27	177	66.3	496	AAW19584	Mouse apoptosis in
28	177	66.3	496	AAW69297	Murine XIAP protei
29	145	54.3	236	AAW61440	Human IAP (an inh
30	145	54.3	236	AAW61440	Human IAP-like pro
31	145	54.3	236	AAW61440	Human IAP-like pro
32	145	54.3	236	AAW61440	Human IAP-like pro
33	145	54.3	236	AAW61440	Human IAP-like pro
34	145	54.3	236	AAW61440	Human IAP-like pro
35	145	54.3	236	AAW61440	Human IAP-like pro
36	145	54.3	236	AAW61440	Human IAP-like pro
37	145	54.3	236	AAW61440	Human IAP-like pro
38	145	54.3	236	AAW61440	Human IAP-like pro
39	145	54.3	236	AAW61440	Human IAP-like pro
40	145	54.3	236	AAW61440	Human IAP-like pro
41	145	54.3	236	AAW61440	Human IAP-like pro
42	145	54.3	236	AAW61440	Human IAP-like pro
43	145	54.3	236	AAW61440	Human IAP-like pro
44	145	54.3	236	AAW61440	Human IAP-like pro
45	145	54.3	236	AAW61440	Human IAP-like pro

ALIGNMENTS

RESULT 1

ID AAW13550 standard; Protein: 46 AA.

AC AAW13550;

DT 22-Jul-1997 (first entry)

XX Human c-IAP2 repeat 2.

DE Human c-IAP2 repeat 2.

KM IAP, inhibitor; apoptosis; RING finger domain; restinosis;

KW myocardial infarction; nephritis; HIV.

XX Homo sapiens.

OS W09706182-A1.

PN 20-FEB-1997.

PD 20-FEB-1997.

XX 06-AUG-1996; 96WO-0512860.

PE 08-DEC-1995; 95US-0569749.

PR 08-AUG-1995; 95US-0512946.

XX (TULAR) TULARIK INC.

PI Goeddel DV, Rothe M;

XX WPI: 1997-154209/14.

DR Nucleic acids encoding cellular inhibitor of apoptosis proteins

PT useful for apoptosis regulation in cells to reduce or increase

P7 apoptosis and for pharmacological screening

XX

PS Claim 3; Page 24; 35pp; English.

XX The human cellular inhibitor of apoptosis proteins (c-IAP1/2 -

CC AAT61590/761591) comprise a series of defined structural domain

CC repeats and/or a RING finger domain; in particular, at least two of

CC a first domain repeat (AAW13547 or AAW13548), a second domain repeat

CC (AAW13549 or AAW13550), and a third domain repeat (AAW13551 or AAW13552)

CC and/or a RING finger domain (AAW13553 or AAW13554), or a consensus

CC sequences derived from these human genes.

CC The nucleic acid is used for recombinant prodn. of human cellular

CC inhibitor of apoptosis protein which modulates apoptosis

CC regulation. The nucleic acids are useful in therapies where

CC increased cell-specific apoptosis is desired, e.g. in restinosis,

CC inflammatory disease states, myocardial infarction, glomerular

CC nephritis, transplant rejection and infectious diseases, e.g. HIV.

CC They can also be used in conditions requiring a reduction in

CC apoptosis.

CC XX

SO Sequence 46 AA;

Query Match 100.0%; Score 267; DB 18; Length 46;

Best Local Similarity 100.0%; Pred. No. 5,4e-27;

Matches 46; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 LAKAGFYITGPDVACFACGCKLSNWEPKDAMSEHLRHPKCP 46

DB 1 lakagfyytgpdvacfacgcklsnwepkdamsehlrhfkpcpf 46

RESULT 2

AAW19747

ID AAW19747 standard; Protein: 604 AA.

XX AAW19747;

DT 16-SEP-1997 (first entry)

XX

DE Human inhibitor of apoptosis protein homologue M1HC.

XX

KW Inhibitor of apoptosis protein; IAP; mammalian IAP homologue; M1HC;

KW degenerative disease; infectious disease; autoimmune disease;

XX cancer; therapy; diagnosis.

XX

OS Homo sapiens.

XX

EH Key Location/Qualifiers

FT Region 29..97

FT /label= BIR

FT Region 169..236

FT /label= BIR

FT Region 255..323

FT /label= BIR

FT Region 556..593

FT /label= RING_finger

XX

PN W09723501-A1.

PD 03-JUL-1997.

XX

PF 20-DEC-1996; 96MO-AU00827.

XX

PR 22-DEC-1995; 95AU-0007275.

XX

PA (AMRA-) AMRAD OPERATIONS PTY LTD.

XX

PI Vaux DL;

XX

DR WPI: 1997-350966/32.

XX

DR N-PSDB: AAT72712.

XX

PT Isolated protein homologues of viral inhibitors of apoptosis - used

PT to modulate apoptosis for treatment of degenerative, infectious or

PT autoimmune diseases and cancer

XX

PS Claim 9; Page 58-62; 136pp; English.

XX

CC Mammalian IAP homologue C (M1HC) (AAW19747) is a human homologue of

CC baculovirus inhibitor of apoptosis protein (IAP). Its amino acid

CC sequence was deduced from a cDNA clone (see also AAT72712) isolated

CC from a human foetal liver cDNA library using primers based on

CC human EST sequences that resembled the BIR repeats of Oryza

CC pseudotsuguta polyhedrosis virus IAP. IAP homologues (see also

CC AAW19745-46 and AAW19748-52) and their derivatives and chemical

CC analogues can be used in methods for modulating apoptosis in animal

CC cells, specifically for treatment, by inhibition, of degenerative

CC and infectious disease or, by promotion, of cancer and autoimmune

CC disease.

CC XX

SO Sequence 604 AA;

Query Match 100.0%; Score 267; DB 18; Length 604;

Best Local Similarity 100.0%; Pred. No. 8,4e-26;

Matches 46; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 LAKAGFYITGPDVACFACGCKLSNWEPKDAMSEHLRHPKCP 46

DB 189 lakagfyytgpdvacfacgcklsnwepkdamsehlrhfkpcpf 234

RESULT 3

AAW13546

ID AAW13546 standard; Protein: 604 AA.

XX AAW13546;

DT 22-JUL-1997 (first entry)

XX

DE Human c-IAP2.

XX

KW IAP; inhibitor; apoptosis; RING finger domain; restinosis;

KW myocardial infarction; nephritis; HIV.

XX

OS Homo sapiens.

XX

OS W09706182-A1.

XX

PD 20-FEB-1997.

XX

PF 06-AUG-1996; 96MO-US12860.

XX

PR 08-DEC-1995; 95US-0569749.

XX

PR 08-AUG-1995; 95US-0512946.

XX

PA (TULA-) TULARIK INC.

XX

PI Goeddel DV, Rothe M;

XX

DR WPI: 1997-154209/14.

XX

DR N-PSDB: AAT61591.

XX

PT Nucleic acids encoding cellular inhibitor of apoptosis proteins -

PT useful for apoptosis regulation in cells to reduce or increase

PT apoptosis and for pharmacological screening

XX

PS Disclosure; Page 21-23; 35pp; English.

XX

CC The human cellular inhibitor of apoptosis proteins (c-IAP1/2 -

CC AAT61590/761591) comprise a series of defined structural domain

CC repeats and/or a RING finger domain; in particular, at least two of

CC a first domain repeat (AAW13547 or AAW13548), a second domain repeat

CC (AAW13549 or AAW13550), and a third domain repeat (AAW13551 or AAW13552)

CC and/or a RING finger domain (AAW13553 or AAW13554), or a consensus

CC sequences derived from these human genes.

CC The nucleic acid is used for recombinant prodn. of human cellular

CC inhibitor of apoptosis protein which modulates apoptosis
CC regulation. The nucleic acids are useful in therapies where
CC increased cell-specific apoptosis is desired, e.g. in restinosis,
CC inflammatory disease states, myocardial infarction, glomerular
CC nephritis, transplant rejection and infectious diseases, e.g. HIV.
CC They can also be used in conditions requiring a reduction in
CC apoptosis.
CC
SQ Sequence 604 AA:

Query Match 100.0%; Score 267; DB 18; Length 604;
Best Local Similarity 100.0%; Pred. No. 8,4e-26;
Matches 46; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 LAKAGFYTGPGDRVACFACGKLSNWEPRKDNAMSEHLRHPKCP 46
DB 189 lakagfyytgpdrvacfaagklsnwepkdnamsehlrhpkcpi 234

RESULT 4

AAV52703 standard; Protein; 604 AA.

AC AAV52703;

DT 26-JAN-2000 (first entry)

DE Human cellular inhibitor of apoptosis-2 protein.

KW Identification: genetic target; gene modulation; human;
KW antisense oligonucleotide; phosphorothioate; target validation;
KW nucleotide sequence-based technology; antisense drug discovery.

OS Homo sapiens.

PN M09953101-A1.

PD 21-OCT-1999.

PF 13-APR-1999; 99WO-US08268.

PR 13-APR-1998; 98US-0081483.

PR 28-APR-1998; 98US-0067638.

PA (ISIS-) ISIS PHARM INC.

PI Cowsett LM, Baker BF, McNeill J, Freiler SM, Sasmor HM, Brooks DG;
PI Ohasi C, Wyatt JR, Borchers AH, Vickers TA;

DR WPI: 1999-620446/53.
DR N-PSDB: AA241005.

PT Identifying compounds which modulate expression of nucleic acids, used
PT to provide compounds having defined physical, chemical or bioactive
PT properties, e.g. antisense activity -

PS Example 20: Page 197-202; 264pp; English.

CC A method has been developed of defining a set of compounds that modulate
CC the expression of a target nucleic acid (tNA) sequence via binding of
CC the compounds with the tNA sequence. The method comprises generating a
CC library of virtual compounds in silico according to defined criteria,
CC and evaluating in silico the binding of the virtual compounds with the
CC tNA according to defined criteria. Also described are: (1) a method of
CC defining a set of oligonucleotides (ONS) that modulate the expression of
CC a tNA sequence via binding of the ONS with the tNA sequence comprising
CC generating a library of virtual compounds in silico according to defined
CC criteria, and evaluating in silico the binding of the virtual ONS with
CC the tNA according to defined criteria; and (2) a method of defining a
CC set of compounds that modulate the expression of a tNA sequence via
CC binding of the compounds with the tNA. The methods can be used for the
CC generation and identification of synthetic compounds having defined

CC physical, chemical or bioactive properties. Information gathered from
CC assays of such compounds is used to identify nucleic acid sequences that
CC are tractable to a variety of nucleotide sequence-based technologies,
CC e.g. antisense drug discovery and target validation. AA240852 to the
CC AA241220, and AAV52701 to AAV52706, represent sequences used in the
CC exemplification of the present invention.
CC
SQ Sequence 604 AA:

Query Match 100.0%; Score 267; DB 20; Length 604;
Best Local Similarity 100.0%; Pred. No. 8,4e-26;
Matches 46; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 LAKAGFYTGPGDRVACFACGKLSNWEPRKDNAMSEHLRHPKCP 46
DB 189 lakagfyytgpdrvacfaagklsnwepkdnamsehlrhpkcpi 234

RESULT 5

AAV33997 standard; Protein; 604 AA.

AC AAV33997;

DT 26-NOV-1999 (first entry)

DE Human cellular inhibitor of apoptosis-2 sequence.

KW Cellular inhibitor of Apoptosis-2; antisense; diagnostic; therapeutic;
KW C-IAP-2; prophylaxis; infection; inflammation; tumor formation.

OS Homo sapiens.

PN US5958771-A.

PD 28-SEP-1999.

PF 03-DEC-1998; 98US-0205144.

PR 03-DEC-1998; 98US-0205144.

PA (ISIS-) ISIS PHARM INC.

PI Bennett CF, Cowsett LM, Ackermann EJ;

DR WPI: 1999-561046/47.
DR N-PSDB: AA222096.

PT Antisense compounds complementary to Cellular Inhibitor of Apoptosis-2
PT useful for e.g. diagnostics, therapeutics, and as research reagents -
PS Example 13: Columns 45-50; 33pp; English.

CC The invention provides antisense compounds of 8-30 nucleotides that
CC inhibit the expression of human Cellular Inhibitor of Apoptosis-2
CC (C-IAP-2). The antisense compounds may be used for diagnostics,
CC therapeutics (for modulating the expression of C-IAP-2), prophylaxis
CC (e.g. to prevent or delay infection, inflammation, or tumor formation),
CC as research reagents (e.g. to distinguish between members of a biological
CC pathway) and in kits. The present sequence represents the human cellular
CC inhibitor of apoptosis-2.

SQ Sequence 604 AA:

Query Match 100.0%; Score 267; DB 20; Length 604;
Best Local Similarity 100.0%; Pred. No. 8,4e-26;
Matches 46; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 LAKAGFYTGPGDRVACFACGKLSNWEPRKDNAMSEHLRHPKCP 46
DB 189 lakagfyytgpdrvacfaagklsnwepkdnamsehlrhpkcpi 234

RESULT 6
AAB50694
ID AAB50694 standard; Protein: 1141 AA.
XX AC
XX AAB50694;
XX DT 19-MAR-2001 (first entry)
XX DE Human API2-MLT chimeric protein sequence.
XX
XX Human: API2-MLT chimera; chimeric; apoptosis inhibitor 2; MTF; API2;
XX mucosa-associated lymphoid tissue lymphoma associated translocation;
XX chromosome 11 region q21-q22.3; chromosome 18 region q21.1-22;
XX molecular characterisation: chromosome translocation; carcinogenesis;
XX fusion protein; malignancy.
XX
XX Chimeric - Homo sapiens.
XX Synthetic.
XX
XX HQ200073500-A1.
XX
XX PD 07-DEC-2000.
XX
XX 26-MAY-2000; 2000QWO-EP04796.
XX
XX 27-MAY-1999; 99EP-0201683.
XX
XX (VLA-) VLAAHS INTERUNIVERSITAIR INST BIOTECHNOG.
XX
XX Baens M, Marynen P, Dierlamm J;
XX
XX WPI: 2001-061556/07.
XX
XX DR N-PSDB: AAC90972.
XX
XX
XX Determining if a tissue sample has a chromosome (11:18) translocation
XX associated with malignancies by amplifying a nucleic acid sample using
XX primers complementary to chromosome 11 region q21-q22.3 and chromosome
XX 18 region q21.1-22 -
XX
XX Claim 12: Fig 5; 47pp: English.
XX
XX
XX The present invention describes a method for determining if a tissue
XX sample comprises a cell with a chromosome (11:18) translocation
XX associated with malignancies such as mucosa-associated lymphoid tissue
XX (MALT) lymphomas. The method comprises subjecting a sample nucleic acid
XX to amplification using primers complementary to sequences which are on
XX chromosome 11 region q21-q22.3 and on chromosome 18 region q21.1-22. The
XX method can be used for determining if a tissue sample or analogue
XX comprises a chromosome (11:18) translocation associated with malignancies
XX such as mucosa-associated lymphoid tissue lymphomas. The nucleic acid or
XX the antibody may be used as a probe for detection for hybridisation to
XX southern blot cell DNAs or for in situ hybridisation of cells, or for
XX determining the presence of complementary DNA. The present sequence
XX represents the specifically claimed chimeric human apoptosis inhibitor 2
XX (API2)/MALT-lymphoma associated translocation (MTF) protein.
XX
XX Sequence 1141 AA:
XX
XX
XX Query Match 100.0%; Score 267; DB 22; Length 1141;
XX Best Local Similarity 100.0%; Pred. No. 1,6e-25;
XX Matches 46; Conservative 0; Mismatches 0; Indels 0; Gaps 0
XX
XX 1 LAAAGPYYIGPGDNACFACGAGKLSMWEKDNAMSEHRLHPKCP 46
XX ||||||||||||||||||||||||||||||||||||||||||||
XX Db 189 Lkkgfyyigpgdnacfacgagklsmwepkdnamselhrhpkcpf 234
XX
XX RESULT 7
XX AA019582 standard; Protein: 604 AA.
XX ID AA019582

[illegible]

AAW69295
ID AAW69295 standard; Protein: 604 AA.
XX
XX AAW69295:
AC
XX
XX 13-NOV-1998 (first entry)
DT
XX
XX Human HIAP-1 protein.
DE
XX
XX Inhibitor of apoptosis protein; apoptosis enhancer; NALP polypeptide;
KW proliferative disease; IAP; therapy; cancer; human; HIAP-1 protein.
XX
XX Homo sapiens.
OS
XX
XX WO9835693-A2.
FN
XX 20-AUG-1998.
PD
XX
XX 13-FEB-1998; 98WO-1800781.
PE
XX
XX 13-FEB-1997; 97US-0800929.
PR
XX
XX (UYOT-) UNIV OTTAWA.
PA
XX
XX Balrd S, Korneluk R, Liston P, Mackenzie AE, Pratt C;
PI Tsang B;
PI
XX
XX WPI: 1998-467164/40.
DR
XX
XX N-PSDB: AAV55039.
DR
XX
XX Inducing apoptosis in proliferative mammalian cells with inhibitor
PT of IAP or NALP polypeptide - also methods for prognosis based on
PT presence of IAP and NALP, specifically applied to cancers involving
PT p53 mutations
PT
XX
XX Disclosure: Fig 2; 147pp; English.
PS
XX
XX This sequence is the human HIAP-1 protein, which is a inhibitor of
CC apoptosis protein (IAP), and can be used in the method of the invention.
CC The method is for enhancing apoptosis in cells from a mammal with
CC proliferative disease by treatment with a compound that inhibits
CC biological activity of an IAP or NALP polypeptide. The inhibitory
CC compounds are used to treat proliferative diseases, specially cancers of
CC ovary, breast, pancreas, lymph nodes, skin, blood, lung, brain, kidney,
CC liver nasopharynx, thyroid, central nervous system, prostate, colon,
CC rectum, cervix or endometrium, particularly to increase their sensitivity
CC to chemotherapeutic agents. High levels of the IAP or NALP proteins are
CC detected in many cancers and are associated with poor prognosis,
CC resistance to chemotherapeutic agents and mutations in p53 (It is
CC suggested that wild-type p53 suppresses transcription of the IAP or NALP
CC genes). Transgenic animals are used for testing the effects of antisense
CC oligonucleotides and for screening for the inhibitors.
CC
XX
XX Sequence 604 AA:
SQ

Query Match 98.9%; Score 264; DB 19; Length 604;
Best Local Similarity 97.8%; Pred. No. 2e-25;
Matches 45; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 1 LAKAGFYTGPGDRVACAFACGGKLSMWEKPKAMSEHRRHPKCP 46
||:|||||
DB 189 laraqfytygpgdrvacafacggklsmwepkdamsehlrhpncpf 234

RESULT 9
ID AAW02925 standard; Protein: 306 AA.
XX
XX AAW02925:
AC
XX
XX 12-SEP-2001 (first entry)
DT
XX

DE Angiotensin converting enzyme (ACEV) splice variant protein #25.
XX
XX Angiotensin converting enzyme splice variant; ACEV; interleukin 6;
KW granulocyte colony stimulating factor receptor; glucagon; hypertrophy;
KW platelet-derived endothelial cell growth factor; cardiovascular disease;
KW cellular tumour antigen p53; cyclin-dependent kinase inhibitor 1C;
KW vasoactive intestinal polypeptide receptor 2; arteriosclerosis; cancer;
KW myocardial infarction; coronary arterial thrombosis; renal disease;
KW diabetic neuropathy; muscular disease; immune disorder; sarcoidosis;
KW multiple sclerosis; immune complex nephritis; deep vein thrombosis;
KW nonrheumatoid pulmonary granulomatous disease; endothelial abnormality;
KW vascular disorder; asbestosis.
XX
XX Homo sapiens.
OS
XX
XX WO200136632-A2.
FN
XX 25-MAY-2001.
PD
XX
XX 17-NOV-2000; 2000MO-1L00766.
PE
XX
XX 17-NOV-1999; 99IL-0132978.
PR
XX
XX 10-DEC-1999; 99IL-0133455.
PR
XX
XX (COMP-) COMPUGEN LTD.
PA
XX
XX Levine Z, David A, Azar I, Khosravi R, Bernstein J;
PI
PI WPI: 2001-336004/35.
DR
XX
XX N-PSDB: AAS06025.
DR
XX
XX Novel alternative splicing variants e.g. variant of angiotensin
PT converting enzyme (ACEV), useful in identifying candidate compounds
PT capable of binding to the variant and to detect anti-variant antibodies
PT
XX
XX Claim 4; Fig 25; 519pp; English.
PS
XX
XX The sequence represents an angiotensin converting enzyme splice variant
CC (ACEV) polypeptide. The polypeptides of the invention include variants of
CC granulocyte colony stimulating factor receptor, glucagon, interleukin 6,
CC platelet-derived endothelial cell growth factor, cyclin-dependent kinase
CC inhibitor 1C, cellular tumour antigen p53, and vasoactive intestinal
CC polypeptide receptor 2. The polypeptides and their associated nucleic
CC acids are useful for identification of variant sequences and detection of
CC candidate compounds capable of binding the molecules. The sequences of
CC the invention can be used in the treatment and diagnosis of various
CC disorders including cardiovascular diseases such as arteriosclerosis,
CC myocardial infarction and coronary arterial thrombosis, renal diseases
CC such as diabetic nephropathy, muscular diseases such as hypertrophy,
CC immune disorders such as immune complex nephritis, multiple sclerosis,
CC cancer, sarcoidosis, nonrheumatoid pulmonary granulomatous diseases such
CC as asbestosis and vascular pathologies involving an endothelial
CC abnormality such as deep vein thrombosis.
CC
XX
XX Sequence 306 AA:
SQ

Query Match 97.0%; Score 259; DB 22; Length 306;
Best Local Similarity 95.7%; Pred. No. 4.3e-25;
Matches 44; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 1 LAKAGFYTGPGDRVACAFACGGKLSMWEKPKAMSEHRRHPKCP 46
||:|||||
DB 204 laraqfytygpgdrvacafacggklsmwepkdamsehlrhpncpf 249

RESULT 10
ID AAW13549 standard; Protein: 46 AA.
XX
XX AAW13549:
AC
XX
XX AAW13549:
DT
XX

```

DF 22-JUL-1997 (first entry)
XX
DE Human c-IAP1 repeat 2.
XX
KW IAP; Inhibitor; apoptosis; RING finger domain; restinosis;
XX myocardial infarction; nephritis; HIV.
XX
OS Homo sapiens.
XX
PN WO9706182-A1.
XX
PD 20-FEB-1997.
XX
PE 06-AUG-1996; 96WO-US12860.
XX
PR 08-DEC-1995; 95US-0569749.
XX
PR 08-AUG-1995; 95US-0512946.
XX
PK (TULSA-) TULARIK INC.
XX
PI Goeddel DV, Rothe M;
XX
DR WPI; 1997-154209/14.
XX
PT Nucleic acids encoding cellular inhibitor of apoptosis proteins
PT useful for apoptosis regulation in cells to reduce or increase
PT apoptosis and for pharmacological screening
XX
PS Claim 3; Page 24; 35pp; English.
XX
CC The human cellular inhibitor of apoptosis proteins (c-IAP1/2 -
CC AAT61590/T61591) comprise a series of defined structural domain
CC repeats and/or a RING finger domain; in particular, at least two of
CC a first domain repeat (AAW13547 or AAW13548), a second domain repeat
CC (AAW13549 or AAW13550), and a third domain repeat (AAW13551 or AAW13552)
CC and/or a RING finger domain (AAW13553 or AAW13554), or a consensus
CC sequence derived from these human genes.
CC The nucleic acid is used for recombinant prodn. of human cellular
CC inhibitor of apoptosis protein which modulates apoptosis
CC regulation. The nucleic acids are useful in therapies where
CC increased cell-specific apoptosis is desired, e.g. in restinosis,
CC inflammatory disease states, myocardial infarction, glomerular
CC nephritis, transplant rejection and infectious diseases, e.g. HIV.
CC They can also be used in conditions requiring a reduction in
CC apoptosis.
XX
SQ Sequence 46 AA:

Query Match          92.9%; Score 248; DB 18; Length 46;
Best Local Similarity 91.3%; Pred. No. 1.4e-24;
Matches 42; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

OY 1 LAKAGFYIIGPGRVACFACGKLSNMEPKDNAMSEHLRHFPKCP 46
   ||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
DB 1 laregfyiyigpgrvactacgklsnwpkddamsehrthfncpl 46

RESULT 11
AAW04583
ID AAW04583 standard; Protein: 438 AA.
XX
AC AAW04583;
XX
DT 07-FEB-1997 (first entry)
XX
DE Human inhibitor of apoptosis gene 1.
XX
KW Inhibitor of apoptosis 1; hiap-1; degenerative disease;
KW rheumatoid arthritis; septic shock; antiviral; trauma; stroke;
KW cell death; oncogenesis; cancer; diagnosis; therapy.
XX
OS Homo sapiens.

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XX
PN WO9635703-A1.
XX
PD 14-NOV-1996.
XX
PF 11-MAY-1995; 95WO-US05922.
XX
PR 11-MAY-1995; 95WO-US05922.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
PI He MW, Hudson PL, Rosen CA;
XX
DR WPI; 1996-518608/51.
XX
DR N-PSDB; AAT43709.
XX
PT Polynucleotide encoding human inhibitor of apoptosis gene 1 - useful
PT for treating degenerative diseases, as antiviral defence mechanism
PT and preventing cell death during trauma and strokes
XX
PS Claim 1; Page 40-41; 53pp; English.
XX
CC Human inhibitor of apoptosis 1 (hiap-1) (AAW04583) is a protein
CC useful for treating degenerative diseases, rheumatoid arthritis,
CC septic shock, as an antiviral defence mechanism, and for
CC preventing cell death during strokes or trauma. Its amino acid
CC sequence was deduced from a cDNA clone (AAT43709) that can be obtd.
CC from human Jurkat cell lines or human osteoclastoma stromal cell
CC lines. Recombinant hiap-1 can be produced in prokaryotic or
CC eukaryotic host cells, or expressed in vivo. It can also be used
CC to screen for modulators of hiap-1 activity.
XX
SQ Sequence 438 AA:

Query Match          92.9%; Score 248; DB 17; Length 438;
Best Local Similarity 91.3%; Pred. No. 1.0e-23;
Matches 42; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

OY 1 LAKAGFYIIGPGRVACFACGKLSNMEPKDNAMSEHLRHFPKCP 46
   ||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
DB 24 laregfyiyigpgrvactacgklsnwpkddamsehrthfncpl 69

RESULT 12
AAW19746
ID AAW19746 standard; Protein: 618 AA.
XX
AC AAW19746;
XX
DT 16-SEP-1997 (first entry)
XX
DE Human inhibitor of apoptosis protein homologue M1HB.
XX
KW Inhibitor of apoptosis protein; IAP; mammalian IAP homologue; M1HB;
KW degenerative disease; infectious disease; autoimmune disease;
KW cancer; therapy; diagnosis.
XX
OS Homo sapiens.
XX
FH Key
FH Region
FT 46..113
FT /label= BIR
FT 184..250
FT /label= BIR
FT 269..337
FT /label= BIR
FT 569..606
FT /label= RING_finger
XX
PN WO9723501-A1.
XX
PD 03-JUL-1997.

```

XX	20-DEC-1996:	96MO-AUD0827.
PF		
XX	22-DEC-1995:	95AU-0007275.
XX		
XX		
PA	(AMRA-)	AMRAD OPERATIONS PTY LTD.
XX		
XX		
PI	Vaux DL:	
DR	WPI: 1997-350966/32.	
DR	N-PSDB: AAT72711.	
PT		
PT	Isolated protein homologues of viral inhibitors of apoptosis - used to modulate apoptosis for treatment of degenerative, infectious or autoimmune diseases and cancer	
XX	Claim 8; Page 51-54; 136pp: English.	
XX		
XX	Mammalian IAP homologue B (MIB) (AAW19746) is a human homologue of baculovirus inhibitor of apoptosis protein (IAP). Its amino acid sequence was deduced from a cDNA clone (see also AAT72711) isolated from a human foetal liver cDNA library using primers based on human EST sequences that resembled the BIR repeats of Oryza pseudotsingua polyhedrosis virus IAP. IAP homologues (see also AAW19745 and AAW19747-52) and their derivatives and chemical analogues can be used in methods for modulating apoptosis in animal cells, specifically for treatment, by inhibition, of degenerative and infectious disease or, by promotion, of cancer and autoimmune disease.	
XX		
SO	Sequence	618 AA:
	Query Match	92.9%; Score 248; DB 18; Length 618;
	Best Local Similarity	91.3%; Pred. No 2,3e-23;
	Matches	42; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
OY	1 LAKAGFTYIGFGDRVACPGAGCLSMWPKDNAMSEHLHPKCF 46	
Db	204 IARAGTYIYIPGDRVACGAGKLSWPKDAMSEHLHPKCF 249	
	RESULT 13	
ID	AAW19583	
XX	AAW19583 standard; protein; 618 AA.	
AC	AAW19583;	
XX		
DT	02-SEP-1997 (first entry)	
XX		
DE	Human apoptosis inhibitor HIAP-2.	
XX		
KW	Apoptosis inhibitor; HIAP-2; HIV; AIDS; neurodegeneration; myelodysplastic syndrome; ischemia; myocardial infarction; stroke; reperfusion injury; toxin-induced liver disease; gene therapy; diagnosis.	
KW		
XX		
OS	Homo sapiens.	
XX		
XX		
FR	Key	Location/Qualifiers
FR	Domain	46..113
FT		/label= BIR-1
FT	Domain	184..250
FT		/label= BIR-2
FT	Domain	269..336
FT		/label= BIR-3
FT	Domain	360..605
FT		/label= Ring_zinc_finger
XX		
XX	WO9706255-A2.	
XX		
PN	20-FEB-1997.	
ED		

Pf	05-AUG-1996;	96MO-IB01022.
Xx	22-DEC-1995;	95US-0576956.
Pr	04-AUG-1995;	95US-0511485.
Xx	(UYOF-) UNIV OTTAWA.	
Pa	Baird S, Korneluk RG, Liston P, Mackenzie AE;	
Pi	WPI; 1997-154262/14.	
Dn	N-PeDB: AAT7083B.	
Dr	Nucleic acid encoding an inhibitor of apoptosis polypeptide - used	
Pt	to inhibit apoptosis in e.g. HIV or AIDS patients, and for detection	
Ft	of susceptibility to apoptotic disease	
Px	Claim 27: Page 75-77; 219pp: English.	
Ss	Human XIAP, HIAP-1 and HIAP-2 and murine M-XIAP, M-HIAP-1 and	
Cc	M-HIAP-2 (AAW19581-86) are a new class of mammalian proteins that	
Cc	are inhibitors of apoptosis (IAP) and which are characterised by	
Cc	the presence of a ring zinc finger domain (see also AAW19587) and at	
Cc	least one BIR (baculovirus IAP repeat) domain (see also AAW19588).	
Cc	The HIAP amino acid sequences were deduced from cDNA clones (AAW70837	
Cc	and AAT70833) from a human liver library. IAP polypeptides can be	
Cc	expressed in host cells (in vitro or in vivo) and used in methods	
Cc	for treating diseases and disorders involving apoptosis, esp. in a	
Cc	human diagnosed as HIV-positive or as having AIDS, a	
Cc	neurodegenerative disease, a myelodysplastic syndrome or an	
Cc	ischemic injury, selected from myocardial infarction, stroke,	
Cc	reperfusion injury, or a toxin-induced liver disease.	
Sq	Sequence 618 AA:	
Qy	Query Match 92.94% Score 248: DB 18: Length 618:	
Db	Best local similarity 91.3%: Pred. No. 2, 3e-23:	
	Matches 42; Conservative 2; Mismatches 2; Indels 0; Gaps 0:	
Oy	1 LAKAGFYIPGGRVACFCGGCKLSNWEKKNAMSEHLHPKPCF 46	
	I::IIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIII	
Db	204 IARGIYIYGPRGVACIACGKAIINWPKDAMSHTIHNPCL 249	
	RESULT 14	
ID	AAW13545 standard; Protein: 618 AA.	
Ac	AAW13545;	
Xx	22-JUL-1997 (First entry)	
Dt	Human c-IAP1.	
Xx	IAP: Inhibitor; apoptosis; RING finger domain; restiposis;	
Kw	myocardial infarction; nephritis; HIV.	
Xx	Homo sapiens.	
Os	WO9706182-A1.	
Pn	20-FEB-1997.	
Pd	06-AUG-1996; 96MO-US12860.	
Pf	08-DEC-1995; 95US-0569749.	
Pr	08-AUG-1995; 95US-0512946.	
Xx	(TULA-) TULARIK INC.	
Pa	Goeddel DV, Roche K;	
Pi	WPI; 1997-154209/14.	

DR N-PSDB: AAT61590.
 XX Nucleic acids encoding cellular inhibitor of apoptosis proteins
 CC useful for apoptosis regulation in cells to reduce or increase
 PT apoptosis and for pharmacological screening
 PS
 XX Disclosure: Page 18-20; 35pp; English.
 CC The human cellular inhibitor of apoptosis proteins (C-IAP1/2 -
 CC AAT61590/761591) comprise a series of defined structural domain
 CC repeats and/or a RING finger domain. In particular, at least two of
 CC a first domain repeat (AAW13547 or AAW13548), a second domain repeat
 CC (AAW13549 or AAW13550), and a third domain repeat (AAW13551 or AAW13552)
 CC and/or a RING finger domain (AAW13553 or AAW13554), or a consensus
 CC sequences derived from these human genes.
 CC The nucleic acid is used for recombinant prodn. of human cellular
 CC inhibitor of apoptosis protein which modulates apoptosis
 CC regulation. The nucleic acids are useful in therapies where
 CC increased cell-specific apoptosis is desired, e.g. in restinosis,
 CC inflammatory disease states, myocardial infarction, glomerular
 CC nephritis, transplant rejection and infectious diseases, e.g. HIV.
 CC They can also be used in conditions requiring a reduction in
 CC apoptosis.
 CC
 SQ Sequence 618 AA:
 Query Match 92.9%; Score 248; DB 18; Length 618;
 Best Local Similarity 91.3%; Pred. No. 2.3e-23;
 Matches 42; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
 OY 1 LAKAGFYIIGPGRVACFACGKLSNMEPRKDNAMSEHLRHPKCP 46
 DB 204 laragfyyigpgrvactacgklsnwepkdamsehrthfncpf 249
 ||:|||||
 RESULT 15
 ID AAW69296 standard; Protein: 618 AA.
 XX
 AC AAW69296;
 DT 13-NOV-1998 (first entry)
 XX
 DE Human HIAP-2 protein.
 XX
 KW Inhibitor of apoptosis protein; apoptosis enhancer; NAIP polypeptide;
 KW proliferative disease; IAP; therapy; cancer; human; HIAP-2 protein.
 XX
 OS Homo sapiens.
 XX
 OS W09835693-A2.
 PN
 XX
 PD 20-AUG-1998.
 XX
 PD 13-FEB-1998; 98WO-IB00781.
 XX
 PE 13-FEB-1997; 97US-0800929.
 XX
 PR 13-FEB-1997; 97US-0800929.
 XX
 PA (UYOT-) UNIV OTTAWA.
 XX
 PI Baird S, Korneluk R, Liston P, Mackenzie AE, Pratt C;
 PI Tsang B;
 XX
 DR WPI: 1998-467164/40.
 DR N-PSDB: AAW55040.
 XX
 XX Inducing apoptosis in proliferative mammalian cells with inhibitor
 PT of IAP or NAIP polypeptide - also methods for prognosis based on
 PT presence of IAP and NAIP, specifically applied to cancers involving
 PT p53 mutations
 XX
 XX Disclosure: Fig 3; 147pp; English.

XX This sequence is the human HIAP-2 protein, which is a inhibitor of
 CC apoptosis protein (IAP), and can be used in the method of the invention.
 CC The method is for enhancing apoptosis in cells from a mammal with
 CC proliferative disease by treatment with a compound that inhibits
 CC biological activity of an IAP or NAIP polypeptide. The inhibitor
 CC compounds are used to treat proliferative diseases, specially cancers of
 CC liver, breast, pancreas, lymph nodes, skin, blood, lung, brain, kidney,
 CC rectum, cervix or endometrium, particularly to increase their sensitivity
 CC to chemotherapeutic agents. High levels of the IAP or NAIP proteins are
 CC detected in many cancers and are associated with poor prognosis.
 CC resistance to chemotherapeutic agents and mutations in p53 (it is
 CC suggested that wild-type p53 suppresses transcription of the IAP or NAIP
 CC genes). Transgenic animals are used for testing the effects of antisense
 CC oligonucleotides and for screening for the inhibitors.
 CC
 SQ Sequence 618 AA:
 Query Match 92.9%; Score 248; DB 19; Length 618;
 Best Local Similarity 91.3%; Pred. No. 2.3e-23;
 Matches 42; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
 OY 1 LAKAGFYIIGPGRVACFACGKLSNMEPRKDNAMSEHLRHPKCP 46
 DB 204 laragfyyigpgrvactacgklsnwepkdamsehrthfncpf 249
 ||:|||||
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Tue Jan 8 08:23:48 2002

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Page 9

